

# Document made available under the Patent Cooperation Treaty (PCT)

International application number: PCT/US05/001297

International filing date: 13 January 2005 (13.01.2005)

Document type: Certified copy of priority document

Document details: Country/Office: US  
Number: 60/545,777  
Filing date: 18 February 2004 (18.02.2004)

Date of receipt at the International Bureau: 18 February 2005 (18.02.2005)

Remark: Priority document submitted or transmitted to the International Bureau in compliance with Rule 17.1(a) or (b)



World Intellectual Property Organization (WIPO) - Geneva, Switzerland  
Organisation Mondiale de la Propriété Intellectuelle (OMPI) - Genève, Suisse

1784211

# THE UNITED STATES OF AMERICA

TO ALL TO WHOM THESE PRESENTS SHALL COME:

UNITED STATES DEPARTMENT OF COMMERCE

United States Patent and Trademark Office

*February 10, 2005*

THIS IS TO CERTIFY THAT ANNEXED HERETO IS A TRUE COPY FROM THE RECORDS OF THE UNITED STATES PATENT AND TRADEMARK OFFICE OF THOSE PAPERS OF THE BELOW IDENTIFIED PATENT APPLICATION THAT MET THE REQUIREMENTS TO BE GRANTED A FILING DATE.

APPLICATION NUMBER: 60/545,777

FILING DATE: *February 18, 2004*

RELATED PCT APPLICATION NUMBER: *PCT/US05/01297*



Certified by

Under Secretary of Commerce  
for Intellectual Property  
and Director of the United States  
Patent and Trademark Office

16569 U.S. PTO  
021804

17631 (AP) PROV

PATENT  
22856 U.S. PTO  
60/54577

**IN THE UNITED STATES PATENT AND TRADEMARK OFFICE**

In re Provisional Application of  
**PATRICK M. HUGHES**

For: METHODS AND COMPOSITIONS FOR THE  
ADMINISTRATION OF PRODRUGS OF PROTON  
PUMP INHIBITORS

**PROVISIONAL APPLICATION FOR PATENT COVER SHEET**

Mail Stop: Provisional Patent Application  
Commissioner for Patents  
P.O. Box 1450  
Alexandria, VA 22313-1450

Dear Sir:

This is a request for filing a PROVISIONAL APPLICATION FOR PATENT  
under 37 CFR 1.53(c).

1. The enclosed PROVISIONAL APPLICATION includes the following:
  - a. Specification (27 pgs);
  - b. Claims (6 pgs);
  - c. Abstract (1 pg);
  - d. Drawings (2 sheets).
  - e. Assignment including cover sheet (4 pgs.)

2. Inventors:

Given Name	Residence
PATRICK M. HUGHES	2 SOMERSET DRIVE ALISO VIEJO, CA 92656

3. Fee Payment

Please charge to Deposit Account No. 01-0885 for the filing fee of \$160.00, as well as for any discrepancies that may occur.

4. Postcard Receipt

Applicants submit herewith a Return/Stamped Postcard.

Respectfully submitted,

Date: February 18, 2004

By: Brent A. Johnson  
Brent A. Johnson, Ph.D.  
Patent Agent  
Registration No. 51,851  
Tel: 714/246-4348  
Fax: 714/246-4249

Please address all future correspondence to:

Brent A. Johnson, Ph.D.  
ALLERGAN, INC.  
2525 Dupont Drive, T2-7H  
Irvine, CA 92612  
Tel: 714/246-4348  
Fax: 714/246-4249

**CERTIFICATE OF EXPRESS MAIL UNDER 37 C.F.R. §1.10**

I hereby certify that this Provisional Patent Application and the documents referred to as enclosed herein are being deposited with the United States Postal Service on February 18, 2004 in an envelope as "Express Mail Post Office To Addressee" mailing label number EV193720795US with sufficient postage for Express Mail addressed to Mail Stop Provisional Patent Application, Commissioner for Patents, P.O. Box 1450, Alexandria, VA 22313-1450.

Date: February 18, 2004

Susan Bartholomew

Name of person mailing paper

Susan Bartholomew

Signature of person mailing paper

**METHODS AND COMPOSITIONS FOR THE ADMINISTRATION OF  
PRODRUGS OF PROTON PUMP INHIBITORS**

By Inventor

PATRICK M. HUGHES

**BACKGROUND OF THE INVENTION**

**Field of the Invention**

**Description of the Related Art**

Benzimidazole derivatives intended for inhibiting gastric acid secretion are disclosed in U.S. Pat. Nos. 4,045,563; 4,255,431; 4,628,098; 4,686,230; 4,758,579; 4,965,269; 5,021,433; 5,430,042 and 5,708,017. Generally speaking, the benzimidazole-type inhibitors of gastric acid secretion are believed to work by undergoing a rearrangement to form a thiophilic species which then covalently binds to gastric H,K-ATPase, the enzyme involved in the final step of proton production in the parietal cells, and thereby inhibits the enzyme. Compounds which inhibit the gastric H,K-ATPase enzyme are generally known in the field as "proton pump inhibitors" (PPI).

Some of the benzimidazole compounds capable of inhibiting the gastric H,K-ATPase enzyme have found substantial use as drugs in human medicine and are known under such names as LANSOPRAZOLE (U.S. Pat. No. 4,628,098), OMEPRAZOLE (U.S. Pat. Nos. 4,255,431 and 5,693,818), ESOMEPRAZOLE (U.S. Pat. No. 6,369,085) PANTOPRAZOLE (U.S. Pat. No. 4,758,579), and RABEPRAZOLE (U.S. Pat. No. 5,045,552). Some of the diseases treated by proton pump inhibitors and specifically by the five above-mentioned drugs include peptic ulcer, heartburn, reflux esophagitis, erosive esophagitis, non-ulcer dyspepsia, infection by *Helicobacter pylori*, alrynitis and asthma.

Whereas the proton pump inhibitor type drugs represent a substantial advance in the field of human and veterinary medicine, they are not totally without shortcomings or disadvantages. For example, it is believed that the short systemic half-life of the drug limits the degree of gastric acid suppression currently achieved. Furthermore, it appears that the short plasma half-life of the drug may contribute to significant gastric pH fluctuations that occur several times a day in patients undergoing PPI therapy. Additionally, PPIs are acid-labile, and in most cases it is necessary to enterically coat the drug in order to prevent the acidic milieu of the stomach from destroying the drug before the drug is absorbed into systemic circulation. Thus, any contribution that might improve the acid stability or plasma half-life of the presently used proton pump inhibitors will be a significant improvement in the art.

As further pertinent background to the present invention, applicants note the concept of prodrugs which is well known in the art. Generally speaking, prodrugs are derivatives of per se drugs, which after administration undergo conversion to the physiologically active species. The conversion may be spontaneous, such as hydrolysis in the physiological environment, or may be enzyme catalyzed. From among the voluminous scientific literature devoted to prodrugs in general, the foregoing examples are cited: Design of Prodrugs (Bundgaard H. ed.) 1985 Elsevier Science Publishers B. V. (Biomedical Division), Chapter 1; Design of Prodrugs: Bioreversible derivatives for various functional groups and chemical entities (Hans Bundgaard); Bundgaard et al. Int. J. of Pharmaceutics 22 (1984) 45-56 (Elsevier); Bundgaard et al. Int. J. of Pharmaceutics 29 (1986) 19-28 (Elsevier); Bundgaard et al. J. Med. Chem. 32 (1989) 2503-2507 Chem. Abstracts 93, 137935y (Bundgaard et al.); Chem. Abstracts 95, 138493f (Bundgaard et al.); Chem. Abstracts 95, 138592n (Bundgaard et al.); Chem. Abstracts 110, 57664p (Alminger et al.); Chem. Abstracts 115, 64029s (Buur et al.); Chem. Abstracts 115, 189582y (Hansen et al.); Chem. Abstracts 117, 14347q (Bundgaard et al.); Chem. Abstracts 117, 55790x (Jensen et al.); and Chem. Abstracts 123, 17593b (Thomsen et al.).

A publication by *Sih., et al.* (Journal of Medicinal Chemistry, 1991, vol. 34, pp 1049-1062), describes N-acyloxyalkyl, N-alkoxycarbonyl, N-

(aminoethyl), and N-alkoxyalkyl derivatives of benzimidazole sulfoxide as prodrugs of proton-pump inhibitors. According to this article these prodrugs exhibited improved chemical stability in the solid state and in aqueous solutions, but had similar activity or less activity than the corresponding parent compounds having a free imidazole N-H group

United States Patent No. 6,093,734 and PCT Publication WO 00109498 (published on February 24, 2000) describe prodrugs of proton pump inhibitors which include a substituted arylsulfonyl moiety attached to one of the benzimidazole nitrogens of proton pump inhibitors having the structure identical with or related to proton pump inhibitor drugs known by the names LANSOPRAZOLE, OMEPRAZOLE, PANTOPRAZOLE and RABEPRAZOLE.

PCT Publication WO 02/30920 describes benzimidazole compounds which are said to have gastric acid secretion inhibitory and anti *H. pylori* effects. PCT Publication WO 02/00166 describes compounds that are said to be nitric oxide (NO) releasing derivatives of proton pump inhibitors of the benzimidazole structure.

U.S. Patent Application having the title "PRODRUGS OF PROTON PUMP INHIBITORS", filed July 15, 2003 by applicants Michael E. Garst, George Sachs, and Jai M. Shin, which has not yet been assigned a serial number, discloses prodrugs of the proton pump inhibitor type drugs having an arylsulfonyl group with an acidic functional group attached, which provided improved solubility in physiological fluids and improved cell penetration.

#### BRIEF DESCRIPTION OF THE INVENTION

Disclosed herein are dosage forms comprising a prodrug of a proton pump inhibitor comprising a biological leaving group bonded to a nitrogen atom of a benzimidazole moiety of said proton pump inhibitor, wherein said dosage form does not comprise a salt of phosphoric acid, and wherein conversion of said prodrug to said proton pump inhibitor depends upon cleavage of a sulfonyl bond.

Also disclosed herein is a method of reducing gastric acid secretion comprising administering to a mammal an effective amount of a sulfonyl prodrug of a proton pump inhibitor in a composition suitable for said administration, provided said composition does not comprise a phosphate buffer.

The use of a sulfonyl prodrug of a proton pump inhibitor for the manufacture of a medicament for the reduction of gastric acid secretion, wherein said medicament does not comprise a phosphate buffer is also disclosed herein.

A pharmaceutical product comprising a composition comprising sulfonamide prodrug of a proton pump inhibitor, and a package for dispensing or storing said prodrug, wherein said composition does not comprise an anionic buffer, is also disclosed herein.

#### Brief Description of the Drawing Figures

Figure 1 is a plot of the % of the original concentration of compound remaining over time. The original concentration of compound 1 was 0.02 mg/mL, and stability was assessed at 25 °C in 1) water, 2) NaCl salt ( $\mu = 0.15$ ), 3) NaCl salt ( $\mu = 0.5$ ), 4) phosphate buffer (pH 7.0,  $\mu = 0.15$ ), and 5) phosphate buffer (pH 7.0,  $\mu = 0.5$ ).

Figure 2 is a log plot of the data of Figure 1.

#### DETAILED DESCRIPTION OF THE INVENTION

While not intending to limit the scope of the invention in any way, or to be bound in any way by theory, we have surprisingly discovered that monovalent, divalent, and trivalent phosphate ions, and/or phosphate buffers significantly destabilize the prodrug compounds disclosed herein. In other words  $\text{PO}_4^{3-}$ ,  $\text{HPO}_4^{2-}$ ,  $\text{H}_2\text{PO}_4^-$ , and/or buffers consisting of combinations of these ions, have an adverse effect upon the stability of prodrugs of proton pump

inhibitors contemplated herein. While not intending to be bound in any way by theory, the aqueous stability of the prodrugs disclosed herein is also believed to be relevant to the stability of solid compositions comprising the prodrugs due to the hygroscopic nature of the compounds.

5       The term “prodrug” has the meaning previously described herein, and in relation to this disclosure refers to a prodrug of a proton pump inhibitor. The term “proton pump inhibitor” also has the meaning previously described herein.

10       The term “dosage form” used in relation to this invention should be interpreted to mean any form of solid or liquid, or combination thereof, which is intended to be administered to a person, including solutions, suspensions, emulsions, and combinations thereof.

15       While not intending to limit the scope of the invention in any way, or to bound in any way by theory, it is believed that phosphate may act as a nucleophile, which attacks the sulfonyl moiety of the prodrug, and thus catalyzes the cleavage of the S-N bond, resulting in the formation of the parent PPI compound. As a result, it is believed that other polyvalent anions may also destabilize the prodrugs disclosed herein. Therefore, certain embodiments relate to dosage forms or compositions which do not comprise a polyvalent anion. The term “polyvalent anion” has the term generally understood by those  
20 of ordinary skill in the art, i.e. a polyvalent anion is an ion having a charge more negative than -1, e.g. -2, -3, -4, etc.

25       While not intending to be bound in any way by theory, it is believed that the sulfonyl group, which is derived from a hard acid, may be more susceptible to attack by hard polyvalent anions, according to the generally known and accepted theory related to the reactivity of hard and soft ions. Additionally, hard ions, being more compact, are less likely to be influenced by steric repulsions in approaching the sulfonyl group, the sulfur atom of which has four ligands. Hardness in many cases may be related to the molecular mass of an ion, as seen by the table below, where the harder ions such as carbonate, phosphate, and sulfate, have lower molecular masses than the softer ions.  
30       Additionally, smaller ions, regardless of hardness are more likely to destabilize

the prodrugs disclosed herein due to the lower susceptibility to unfavorable steric interactions with the sulfonyl group.

Ion	Molecular Mass (-1 ion)
Carbonate	61
Phosphate	97
Sulfate	97
Malonate	103
Succinate	117
Tartrate	149
Citrate	191

5           Thus, certain embodiments relate to the molecular mass of an ion. The term “molecular mass” has the meaning generally understood in the art, that is, it is the sum of the atomic masses of all individual atoms in a molecule or ion. For the purposes of this disclosure, the term molecular mass is also applicable to ions consisting of only one atom. In one embodiment the prodrug is in a dosage  
10 form or a composition which does not comprise a polyvalent anion having a molecular mass of 100 or less. In another embodiment the prodrug is in a dosage form or a composition which does not comprise a polyvalent anion having a molecular mass of 102 or less. In another embodiment the prodrug is in a dosage form or a composition which does not comprise a polyvalent anion  
15 having a molecular mass of 110 or less. In another embodiment the prodrug is in a dosage form or a composition which does not comprise a polyvalent anion having a molecular mass of 120 or less.

Certain embodiments also relate to the solubility of an ion. While not intending to be bound in any way by theory, it is believed that a more soluble  
20 anion is more likely to contribute to the instability of the prodrug since a higher concentration of the anion can be present in an aqueous environment, thus increasing the kinetic instability of the compound. The “solubility” as used herein in relation to the concentration of the ion is the concentration of the ion in water when the ion is saturated. Since solubility is dependent upon other  
25 components present in a composition, for the purposes of the claim elements, the “solubility” is the concentration of the anion in water when the entire

composition in which the anion is present is intimately contacted with water, and the water is saturated with the anion.

In one embodiment the prodrug is in a dosage form or a composition which does not comprise a polyvalent anion having an aqueous solubility of 0.2 M or greater. In another embodiment the prodrug is in a dosage form or a composition which does not comprise a polyvalent anion having an aqueous solubility of 0.15 M or greater. In another embodiment the prodrug is in a dosage form or a composition which does not comprise a polyvalent anion having an aqueous solubility of 0.1 M or greater. In another embodiment the prodrug is administered in a dosage form or a composition which does not comprise a polyvalent anion having an aqueous solubility of 0.02 M or greater. In another embodiment the prodrug is in a dosage form or a composition which does not comprise a polyvalent anion having an aqueous solubility of 0.015 M or greater. In another embodiment the prodrug is in a dosage form or a composition which does not comprise a polyvalent anion having an aqueous solubility of 0.01 M or greater.

In one embodiment the prodrug is in a dosage form or a composition which does not comprise an anion having an aqueous solubility of 0.1 M or greater and a molecular mass of 110 or less. In another embodiment the prodrug is in a dosage form or a composition which does not comprise an anion having an aqueous solubility of 0.01 M or greater and a molecular mass of 110 or less. In another embodiment the prodrug is in a dosage form or a composition which does not comprise an anion having an aqueous solubility of 0.15 M or greater and a molecular mass of 120 or less. In another embodiment the prodrug is in a dosage form or a composition which does not comprise an anion having an aqueous solubility of 0.015 M or greater and a molecular mass of 120 or less.

In one embodiment the prodrug is in a dosage form or a composition which does not comprise an anionic buffer. The term "buffer" as used herein should be construed to have a narrow meaning according to that which is generally understood in the art. That is, not only should the "buffer" have one or more of the required components which make it a buffer, but the buffer

should be at such a concentration as to be effective in maintaining the pH at the desired value. A phosphate buffer is a combination of phosphoric acid and its salts in a ratio and at an effective concentration, such that the pH is maintained at its desired value for as long as necessary. The desired value of the pH and  
5 the amount of time that the pH must be maintained at that value are dependent upon the composition or dosage form in which the drug is present. Such a determination can be readily made by a person of ordinary skill in the art.

Another embodiment comprises a dosage form or composition comprising a prodrug and a buffer which is not anionic. Buffers which are not  
10 anionic include zwitterionic buffers comprising amino acids such as glycine, or other zwitterionic species such as betaines, and cationic buffers including amines such as triethanolamine or diethanolamine and their salts.

In one embodiment the prodrug is in a dosage form or a composition which does not comprise more than 0.1 moles of a polyvalent anion for every 1  
15 mole of said prodrug, wherein the polyvalent anion has an aqueous solubility of 0.1 M or greater. In another embodiment the prodrug is in a dosage form or a composition which does not comprise more than 0.05 moles of a polyvalent anion for every 1 mole of said prodrug, wherein said polyvalent anion has an aqueous solubility of 0.15 M or greater.

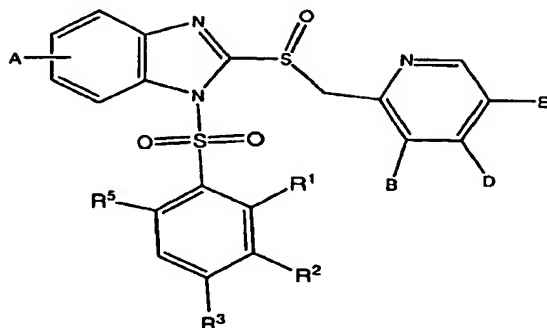
20 The term "biological leaving group" as used herein refers to a moiety which is cleaved from the remainder of the molecule in the body of a mammal such that the remainder of the molecule is a proton pump inhibitor, or is readily converted to a proton pump inhibitor by a process such a protonation; deprotonation; quenching of an unstable intermediate such as a radical, radical  
25 ion, carbocation, carbene, or nitrene; tautomerization; or a similar process. In one embodiment the, biological leaving group comprises a sulfonyl group, where the sulfur atom is directly bonded to the nitrogen atom of the benzimidazole moiety. A "sulfonyl" moiety or group is defined herein as a moiety comprising an SO<sub>2</sub> group, where a sulfur atom is directly covalently  
30 bonded to two oxygen atoms. A "sulfonyl bond" is a bond between the sulfur of the sulfonyl group and another atom. In another embodiment, the biological leaving group comprises a sulfonyl group and an aromatic ring, wherein the

sulfur atom is directly bonded to the nitrogen atom of the benzimidazole moiety. The term "aromatic ring" has the broadest meaning generally understood in the art. In another embodiment, the biological leaving group comprises a phenylsulfonyl group, wherein the sulfur atom is directly bonded to the nitrogen atom of the benzimidazole moiety. The term "phenylsulfonyl" moiety should be broadly interpreted to mean any moiety where the sulfur of the SO<sub>2</sub> group is directly covalently bonded to a carbon that is part of a phenyl ring. The term "phenyl ring" should be broadly understood to mean any ring comprising six carbon atoms having three conjugated double bonds. Thus, a phenylsulfonyl moiety could be monosubstituted, meaning that the sulfonyl group is the only group directly attached to the phenyl ring, or the phenylsulfonyl moiety could have from 1 to 5 additional substituents which are not a hydrogen atom, and are directly attached to a carbon of the phenyl ring.

While not intending to limit the scope of the invention in any way, in many situations one might choose a prodrug which would be converted after administration into one of the widely used and well tested commercially available proton pump inhibitors (PPI) such as lansoprazole, esomeprazole, omeprazole, pantoprazole, and rabeprazole. In situations where one of the commercially available PPIs is used, one may want to consider circumstances related to the individual to which the prodrug is administered in making decisions related to the choice of the compound used. For example, if the person to which the prodrug is being administered is known to respond well to omeprazole, then one may consider using a prodrug of omeprazole as disclosed herein. In another situation, a person may have a history of being effectively treated by lansoprazole, in which case one may consider using a prodrug of lansoprazole as disclosed herein. The specific disclosure related to the proton pump inhibitor is given herein merely to provide guidance and direction to one practicing the disclosure herein, and is not intended to limit the overall scope of the invention in any way.

Certain embodiments relate to particular structures, which are useful as prodrugs.

One embodiment comprises



or a pharmaceutically acceptable salt thereof  
wherein

A is H, OCH<sub>3</sub>, or OCHF<sub>2</sub>;

5 B is CH<sub>3</sub> or OCH<sub>3</sub>;

D is OCH<sub>3</sub>, OCH<sub>2</sub>CF<sub>3</sub>, or O(CH<sub>2</sub>)<sub>3</sub>OCH<sub>3</sub>;

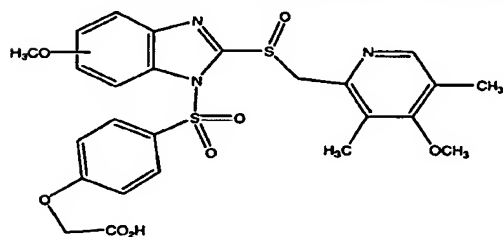
E is H or CH<sub>3</sub>;

R<sup>1</sup>, R<sup>2</sup>, R<sup>3</sup>, and R<sup>5</sup> are independently H, CH<sub>3</sub>, CO<sub>2</sub>H, CH<sub>2</sub>CO<sub>2</sub>H, (CH<sub>2</sub>)<sub>2</sub>CO<sub>2</sub>H,  
CH(CH<sub>3</sub>)<sub>2</sub>, OCH<sub>2</sub>C(CH<sub>3</sub>)<sub>2</sub>CO<sub>2</sub>H, OCH<sub>2</sub>CO<sub>2</sub>CH<sub>3</sub>, OCH<sub>2</sub>CO<sub>2</sub>H, OCH<sub>2</sub>CO<sub>2</sub>NH<sub>2</sub>,

10 OCH<sub>2</sub>CONH<sub>2</sub>(CH<sub>2</sub>)<sub>5</sub>CO<sub>2</sub>CH<sub>3</sub>, or OCH<sub>3</sub>.

In another embodiment related to the one just described, R<sup>1</sup>, R<sup>2</sup>, R<sup>3</sup>, and  
R<sup>5</sup> are independently H, CH<sub>3</sub>, CO<sub>2</sub>H, CH<sub>2</sub>CO<sub>2</sub>H, (CH<sub>2</sub>)<sub>2</sub>CO<sub>2</sub>H, OCH<sub>2</sub>CO<sub>2</sub>CH<sub>3</sub>,  
OCH<sub>2</sub>CO<sub>2</sub>H, OCH<sub>2</sub>CONH<sub>2</sub>(CH<sub>2</sub>)<sub>5</sub>CO<sub>2</sub>CH<sub>3</sub>, or OCH<sub>3</sub>.

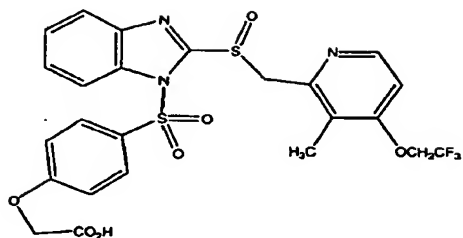
In certain embodiments, the prodrug has a structure comprising



15

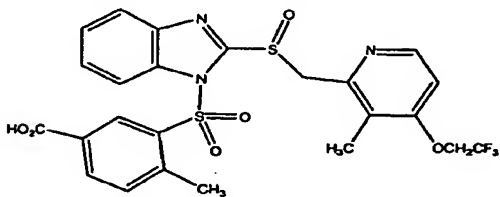
or a pharmaceutically acceptable salt thereof.

Other embodiments comprise



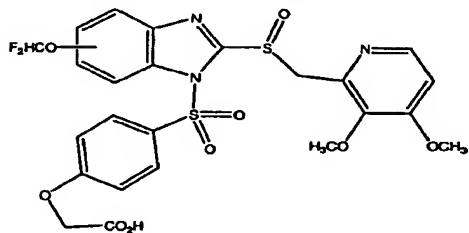
or a pharmaceutically acceptable salt thereof.

Other embodiments comprise



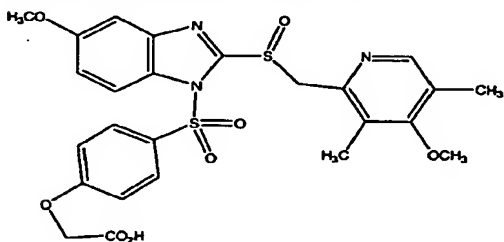
5 or a pharmaceutically acceptable salt thereof.

Other embodiments comprise



or a pharmaceutically acceptable salt thereof.

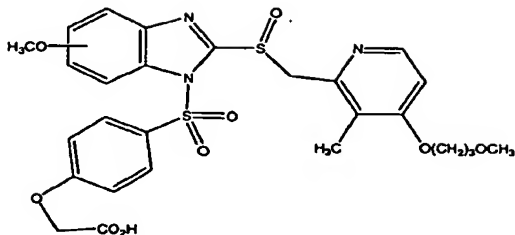
Other embodiments comprise



10

or a pharmaceutically acceptable salt thereof.

Other embodiments comprise



or a pharmaceutically acceptable salt thereof.

A "pharmaceutically acceptable salt" is any salt that retains the activity of the parent compound and does not impart any deleterious or untoward effect on the subject to which it is administered and in the context in which it is administered. Pharmaceutically acceptable salts may be derived from organic or inorganic bases. The salt may be a mono or polyvalent ion. Of particular interest are the inorganic ions, lithium, sodium, potassium, calcium, and magnesium. Organic salts may be made with amines, particularly ammonium salts such as mono-, di- and trialkyl amines or ethanol amines. Salts may also be formed with caffeine, tromethamine and similar molecules. Hydrochloric acid or some other pharmaceutically acceptable acid may form a salt with a compound that includes a basic group, such as an amine or a pyridine ring.

The prodrugs of the present invention can be prepared by the methods described in the following U.S. Patent documents, all of which are expressly incorporated by reference herein: U.S. Pat. No. 6,093,734; U.S. Pat. App. No. 09/783,807, filed February 14, 2001; the U.S. Pat. App. having the title "PRODRUGS OF PROTON PUMP INHIBITORS", filed July 15, 2003 by applicants Michael E. Garst, George Sachs, and Jai M. Shin, which has not yet been assigned a serial number; and the U.S. Pat. App. having the title "PROCESS FOR PREPARING ISOMERICALLY PURE PRODRUGS OF PROTON PUMP INHIBITORS", filed July 15, 2003 by applicants Michael E. Garst, Lloyd J. Dolby, Shervin Esfandiari, Vivian R. Mackenzie, Alfred A. Avey, Jr., David C. Muchmore, Geoffrey K. Cooper, and Thomas C. Malone, which has not yet been assigned a serial number. However, these methods are only given to provide guidance, and are not meant to limit the scope of the invention in any way. One of ordinary skill in the art will recognize that there

are many ways in which the prodrugs of the present invention can be prepared without departing from the spirit and scope of the present invention.

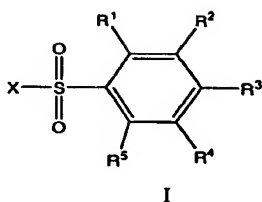
Those skilled in the art will readily understand that for oral administration the compounds of the invention are admixed with  
5 pharmaceutically acceptable excipients which per se are well known in the art. Specifically, a drug to be administered systemically, it may be confectioned as a powder, pill, tablet or the like, or as a syrup or elixir suitable for oral administration. Description of the substances normally used to prepare tablets,  
10 powders, pills, syrups and elixirs can be found in several books and treatise well known in the art, for example in Remington's Pharmaceutical Science, Edition 17, Mack Publishing Company, Easton, Pa.

Parenteral administration is generally characterized by injection. Injectables can be prepared in conventional forms, either as liquid solutions or suspensions, solid forms suitable for dissolving or suspending in liquid prior to  
15 injection, or as emulsions. Descriptions of substances and methods normally used to prepare formulations for parenteral administration can be found in several treatises and books well known in the art such as, Handbook On Injectable Drugs (11th edition), edited by Lawrence A. Trissel, (Chicago: Login Brothers Book Company; January 15, 2001).

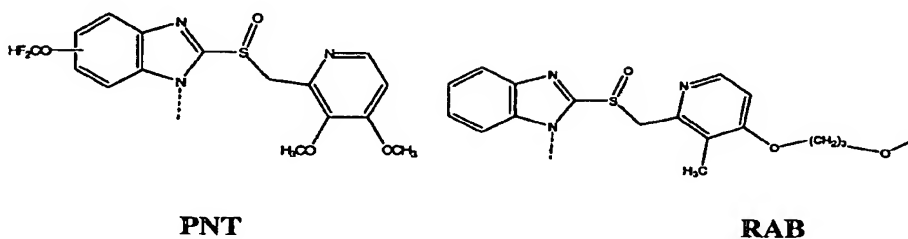
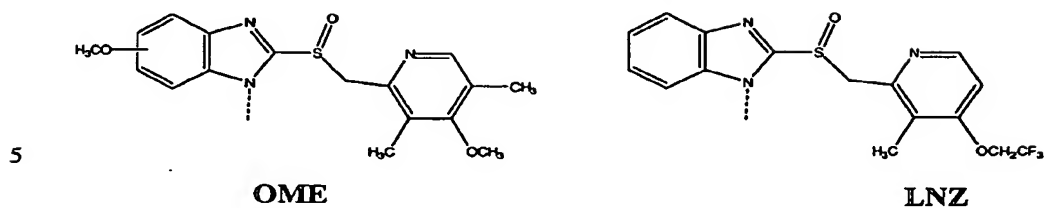
20 The following examples provide guidance and direction in making and using the invention. However, they are not to be interpreted as limiting the scope of the invention in any way.

#### Example 1

25 Compounds specifically contemplated in relation to embodiments disclosed herein are presented in Table 1 below. The generic structure, I, is shown as a combination of a proton pump inhibitor (X) and a sulfonyl-bearing moiety which is attached to the proton pump inhibitor to form the prodrug  
30 according to the formula below. The identity of each group represented by R<sup>1</sup>-R<sup>5</sup> is shown in the table.



The different possibilities for X are shown below.



10

Table 1

Compound	X	R <sup>1</sup>	R <sup>2</sup>	R <sup>3</sup>	R <sup>4</sup>	R <sup>5</sup>
1	OME	H	H	OCH <sub>2</sub> CO <sub>2</sub> H	H	H
2	OME	CH <sub>3</sub>	H	OCH <sub>2</sub> CO <sub>2</sub> H	H	CH <sub>3</sub>
3	OME	H	H	OCH <sub>2</sub> C(CH <sub>3</sub> ) <sub>2</sub> CO <sub>2</sub> H	H	H
4	OME	CH <sub>3</sub>	H	OCH <sub>2</sub> C(CH <sub>3</sub> ) <sub>2</sub> CO <sub>2</sub> H	H	CH <sub>3</sub>
5	OME	H	H	CH <sub>2</sub> CO <sub>2</sub> H	H	H
6	OME	H	CO <sub>2</sub> H	H	H	H
7	LNZ	H	CO <sub>2</sub> H	H	H	H
8	LNZ	H	CO <sub>2</sub> H	OCH <sub>3</sub>	H	H
9	LNZ	H	H	CH <sub>2</sub> CO <sub>2</sub> H	H	H
10	LNZ	H	H	OCH <sub>2</sub> CO <sub>2</sub> H	H	H
11	LNZ	H	H	OCH <sub>2</sub> C(CH <sub>3</sub> ) <sub>2</sub> CO <sub>2</sub> H	H	H
12	LNZ	H	CH <sub>2</sub> CO <sub>2</sub> H	CH <sub>2</sub> CO <sub>2</sub> H	H	H
13	LNZ	H	CO <sub>2</sub> H	H	H	CH <sub>3</sub>
14	LNZ	H	CO <sub>2</sub> H	H	H	OCH <sub>3</sub>
15	LNZ	CH(CH <sub>3</sub> ) <sub>2</sub>	H	CH <sub>2</sub> CO <sub>2</sub> H	H	H
16	LNZ	H	OCH <sub>2</sub> CO <sub>2</sub> H	CO <sub>2</sub> H	H	H

17	LNZ	CH(CH <sub>3</sub> ) <sub>2</sub>	H	OCH <sub>2</sub> CO <sub>2</sub> H	H	CH <sub>3</sub>
18	LNZ	H	H	CO <sub>2</sub> H	H	H
19	LNZ	H	(CH <sub>2</sub> ) <sub>2</sub> CO <sub>2</sub> H	CH <sub>3</sub>	H	H
20	OME	H	H	OCH <sub>2</sub> CO <sub>2</sub> CH <sub>3</sub>	H	H
21	OME	H	H	OCH <sub>2</sub> CO <sub>2</sub> NH <sub>2</sub>	H	H
22	OME	H	CO <sub>2</sub> H	CO <sub>2</sub> H	H	H
23	OME	H	CO <sub>2</sub> H	OCH <sub>2</sub> CO <sub>2</sub> H	H	H
24	OME	H	OCH <sub>2</sub> CO <sub>2</sub> H	OCH <sub>2</sub> CO <sub>2</sub> H	H	H
25	OME	OCH <sub>3</sub>	H	CO <sub>2</sub> H	H	H
26	OME	H		CO <sub>2</sub> H	H	H
27	OME	H	CO <sub>2</sub> H	H	H	CH <sub>3</sub>
28	PNT	H	H	OCH <sub>2</sub> CO <sub>2</sub> H	H	H
29	PNT	H	CO <sub>2</sub> H	H	H	CH <sub>3</sub>
30	RAB	H	CO <sub>2</sub> H	H	H	H
31	RAB	H	CO <sub>2</sub> H	H	H	CH <sub>3</sub>
32	RAB	CH <sub>3</sub>	H	OCH <sub>2</sub> CO <sub>2</sub> H	H	CH <sub>3</sub>
33	RAB	H	H	CO <sub>2</sub> H	H	H
34	LNZ	CH <sub>3</sub>	H	OCH <sub>2</sub> CO <sub>2</sub> H	H	CH <sub>3</sub>
35	LNZ	H	OCH <sub>2</sub> CO <sub>2</sub> H	OCH <sub>2</sub> CO <sub>2</sub> H	H	H
36	LNZ	H	H	CO <sub>2</sub> H	H	H
37	LNZ	CH <sub>3</sub>	H	CO <sub>2</sub> H	H	H
38	LNZ	H	(CH <sub>2</sub> ) <sub>2</sub> CO <sub>2</sub> H	OCH <sub>3</sub>	H	H
39	OME	CH <sub>3</sub>	H	OCH <sub>2</sub> CONH <sub>2</sub> (CH <sub>2</sub> ) <sub>5</sub> CO <sub>2</sub> CH <sub>3</sub>	H	CH <sub>3</sub>
40	OME	H	H	OCH <sub>2</sub> CONH <sub>2</sub> (CH <sub>2</sub> ) <sub>5</sub> CO <sub>2</sub> CH <sub>3</sub>	H	H
41	OME	H	H	(CH <sub>2</sub> ) <sub>2</sub> CO <sub>2</sub> H	H	H
42	OME	H	(CH <sub>2</sub> ) <sub>2</sub> CO <sub>2</sub> H	OCH <sub>3</sub>	H	H

These compounds have been prepared according to procedures described the U.S. Pat. App. having the title "PRODRUGS OF PROTON PUMP

- 5 INHIBITORS", filed July 15, 2003 by applicants Michael E. Garst, George Sachs, and Jai M. Shin, which has not yet been assigned a serial number; and the U.S. Pat. App. having the title "PROCESS FOR PREPARING ISOMERICALLY PURE PRODRUGS OF PROTON PUMP INHIBITORS ", filed July 15, 2003 by applicants Michael E. Garst, Lloyd J. Dolby, Shervin
- 10 Esfandiari, Vivian R. Mackenzie, Alfred A. Avey, Jr., David C. Muchmore, Geoffrey K. Cooper, and Thomas C. Malone, which has not yet been assigned a serial number, incorporated by reference previously herein. These aforementioned patent documents, as well as the provisional U.S. Patent Application No. 513880, filed on October 22, 2003 by applicants Jie Shen,

Devin F. Welty, and Diane D. Tang-Liu, incorporated herein by reference, demonstrate that compounds 1-42 decompose in vivo to form proton pump inhibitors.

5

### Example 2

The physicochemical properties of compound 1 were analyzed. Compound 1 was found to be hygroscopic, in that 9% weight gain was observed for the compound after 14 days of storage at 25 °C at 75% relative humidity.

10

Table 2A. Stability Profile of Compound 1 at 25 °C in Buffered Aqueous Solutions

pH	Buffer Composition	Half-life ( $t_{1/2}$ ) hours	Shelf life ( $t_{90\%}$ ) hours	Degradation Rate Constant (k) 1/hours
1	0.1 M HCl	3.6	0.5	0.194
3	Citric Acid (0.1 M)/ Na <sub>2</sub> HPO <sub>4</sub> (0.2 M)	78.0	11.9	0.009
5	Citric Acid (0.1 M) /Na <sub>2</sub> HPO <sub>4</sub> (0.2 M)	89.2	13.6	0.008
7	sodium phosphate (0.1 - 0.2 M)	286.8	43.6	0.002
7.4	sodium phosphate (0.1 - 0.2 M)	291.2	44.3	0.002
9	sodium phosphate (0.1 - 0.2 M)	23.0	3.5	0.030
10	sodium phosphate (0.1 - 0.2 M)	2.3	0.4	0.298
	No buffer	2863.6	435.4	0.0002

The aqueous stability data of compound 1 is presented in Table 2B. These results show that, the half-life ( $t_{1/2}$ ), the shelf-life ( $t_{90\%}$ ), and the rate constant for degradation (k) for compound 1 are significantly improved at the pH values of 7 and 7.4 relative to the other pH values studied. While not intending to be bound in any way by theory, the fact that compound 1 becomes less stable in both acidic and basic environments, points to both acid and base-catalyzed degradation of these compounds. The base-catalyzed degradation is unexpected because the commercial proton pump inhibitors are stabilized in

15

20

aqueous solutions by adjusting the solution to high pH. In fact, while not intending to be bound or limited in any way by theory, compound 1 appears to be more susceptible to base-catalyzed degradation than acid-catalyzed degradation, since its half-life is longer at pH 5, where the  $H^+$  concentration is  $10^{-5}$  M than its half-life is at pH 9, where the  $OH^-$  concentration is  $10^{-5}$  M. Similarly, compound 1 is less stable at pH 10, where the  $OH^-$  concentration is  $10^{-4}$  M than it is at pH 1, where the  $H^+$  concentration is 0.1 M. While not intending to be bound or limited in any way by theory, these results unexpectedly show that the optimum pH for the compounds disclosed herein is around neutral, and that formulation of aqueous dosage forms of near neutral pH should greatly improve the stability of the prodrugs, thus improving shelf-life and facilitating formulation.

While not intending to be bound or limited in any way by theory, based upon the fact that the stability of compound 1 is essentially unchanged from pH 7 to pH 7.4, and based upon the other data presented in Table 2A, it is reasonable to believe that these compounds should be most stable when the pH is from about pH 6.5 to about 8.

Additionally, these results demonstrate that the prodrugs are significantly more stable in neutral aqueous solutions than the proton pump inhibitors. The stability of omeprazole and other proton pump inhibitors have been reported (Kromer et al., "Differences in pH-Dependent Activation Rates of Substituted Benzimidazoles and Biological in vitro Correlates", Pharmacology 1998; 56:57-70; and Ekpe et al, "Effect of Various Salts on the Stability of Lansoprazole, Omeprazole, and Pantoprazole as Determined by High Performance Liquid Chromatography", Drug Development and Industrial Pharmacy, 25(9), 1057-1065 (1999)), and while the stability is somewhat buffer dependent, typical half-lives for omeprazole are about 40 hours at pH 7, which is nearly an order of magnitude shorter than the prodrug half-life presented in Table 2A. While not intending to be bound in any way by theory, or to limit the scope of the invention in any way, these results suggest that the compounds of disclosed herein can be injected at a more neutral pH than is currently possible with the currently available proton pump inhibitors. This should allow bolus

injection of the compounds disclosed herein as opposed to the slow infusion of the drug currently in practice because the present compositions will not have the irritation associated with the high pH traditionally used with proton pump inhibitors. Additionally, while not intending to be bound in any way by theory, or to limit the invention in any way, these results also demonstrate that the aqueous solution can be stored for a longer period of time prior to administration, and that the solid will be easier to handle, because moisture is less likely to destabilize the active compound.

Surprisingly, we found that the unbuffered prodrug had a half-life that was about an order of magnitude longer than the buffered prodrug. This finding was investigated in detail, and the results are presented in the next example.

### Example 3

The stability of compound 1 at a concentration of 0.02 mg/mL in water at 25 °C was assessed in 1) water, 2) NaCl salt ( $\mu = 0.15$ ), 3) NaCl salt ( $\mu = 0.5$ ), 4) phosphate buffer (pH 7.0,  $\mu = 0.15$ ), and 5) phosphate buffer (pH 7.0,  $\mu = 0.5$ ). For the buffer solutions, the ionic strength ( $\mu$ ) was adjusted using sodium chloride, and the buffer concentration of the two solutions was equal (0.1 M). The amount of remaining compound 1 is presented as the % of the original concentration of 0.02 mg/mL for each sample in Table 3a and in Figure 1. These results show that beyond four days, the stability of the prodrug in the corresponding environment decreased in the following order: water > NaCl salt >> phosphate buffer. The results of the early measurements are anomalous, and suggest an impurity in the sample that may have affected the stability before the impurity was consumed. Figure 2, which is a log plot of the remaining sample, clearly shows a first order decay of the sample from 3-29 days, supporting the hypothesis that the decay of the sample of the first three days are anomalous. The half lives of each sample during this time period were determined, and are presented in Table 3b.

Table 3a

Sampling schedule (Days)	Compound 1 % Remaining				
	Water (pH 7.2)	NaCL, u=0.15 (pH 6.6)	NaCL, u=0.5 (pH 6.2)	Phosphate buffer, u=0.15 (pH 7.0)	Phosphate buffer, u=0.5 (pH 7.0)
1	92.212	84.134	87.598	96.294	96.709
3	86.085	76.960	80.750	90.597	90.087
4	86.410	77.037	78.348	87.434	85.994
7	84.569	75.398	76.513	80.995	76.165
9	84.452	71.861	74.482	76.176	70.768
11	86.930	73.763	74.312	74.520	67.826
13	83.661	72.390	71.691	68.020	n/a
15	80.913	67.858	68.167	62.389	52.494
20	78.768	64.953	65.173	54.360	44.057
24	79.915	64.848	65.753	50.246	39.181
29	78.412	62.731	62.867	43.321	32.626

While not intending to be bound in any way by theory, the fact that the prodrug has a significantly shorter half life and shelf life, and faster decay rate in the phosphate buffer than it had in water at a nearly identical pH demonstrates that phosphate has a destabilizing effect upon the prodrug. While not intending to be bound in any way by theory, it also appears that the presence of other ions may have some adverse effect upon the stability of these compounds, although it is significantly less than that of the phosphate buffer. However, this contribution may simply be a product of the lower pH of the samples.

Decomposition of the prodrug in this and the previous example gave the parent proton pump inhibitor.

Table 3b

Compound 1 sample	Half Life (t <sub>1/2</sub> ) days	Shelf Life (t 90%) days	Rate Constant (k) 1/days
water, pH 7.2	161.163	24.502	0.0043
NaCl, u=0.15, pH 6.6	80.581	12.251	0.0086
NaCl, u=0.5, pH 6.6	75.326	11.452	0.0092

Na phosphate buffer, u=0.15, pH 7.0	24.063	3.658	0.0288
Na phosphate buffer, u=0.5, pH 7.0	17.589	2.674	0.0394

### Example 3

Capsules are prepared according to well-known commercial processes  
5 using the composition shown in Table 3.

Table 3

Component	Amount (mg)
Compound 1	20
Lactose	200
Magnesium Stearate	3

### Example 4

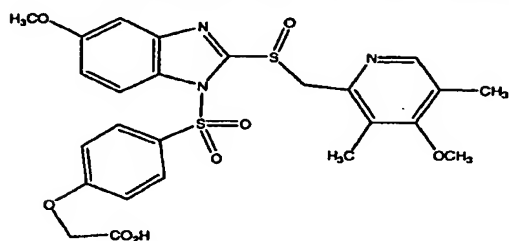
15

The capsule prepared according to example 3 is orally administered daily to a person suffering from heartburn. Relief of pain begins to occur within about 1 day, and continues as long as the person takes the dosage form.

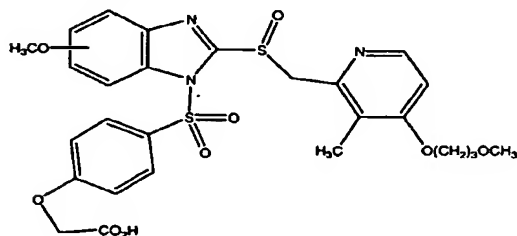
### CLAIMS

What is claimed is:

1. A dosage form comprising  
5 a prodrug of a proton pump inhibitor comprising a biological leaving group bonded to a nitrogen atom of a benzimidazole moiety of said proton pump inhibitor,  
wherein said dosage form does not comprise a salt of phosphoric acid,  
and wherein conversion of said prodrug to said proton pump inhibitor depends  
10 upon cleavage of a sulfonyl bond.
2. The dosage form of claim 1 wherein said proton pump inhibitor is selected from the group consisting of omeprazole, esomeprazole, lansoprazole, pantoprazole, and rabeprazole.
3. The dosage form of claim 1 wherein the proton pump inhibitor is  
15 omeprazole.
4. The dosage form of claim 1 wherein the biological leaving group comprises an phenyl ring.
5. The dosage form of claim 1 comprising



- 20 or a pharmaceutically acceptable salt thereof.
6. The dosage form of claim 1 comprising

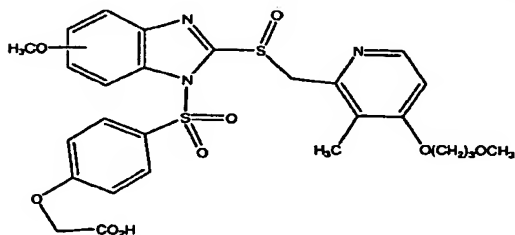


or a pharmaceutically acceptable salt thereof.

7. The dosage form of claim 1 which does not comprise a polyvalent anion having a molecular mass of 100 or less.
8. The dosage form of claim 1 which does not comprise a buffer.
9. The dosage form of claim 1 which does not comprise more than 0.1  
5 moles of a polyvalent anion for every 1 mole of said prodrug, wherein said polyvalent anion has an aqueous solubility of 0.1 M or greater.
10. The dosage form of claim 1 which does not comprise a polyvalent anion having an aqueous solubility of 0.1 M or greater.
11. The dosage form of claim 1 which does not comprise a polyvalent anion  
10 having an aqueous solubility of 0.01 M or greater.
12. The dosage form of claim 6 which does not comprise an anion having an aqueous solubility of 0.1 M or greater and a molecular mass of 110 or less.
13. The dosage form of claim 5 which does not comprise an anion having an aqueous solubility of 0.01 M or greater and a molecular mass of 110 or less.
- 15 14. The dosage form of claim 1 which is a solid.
15. The dosage form of claim 1 which is a liquid.
16. A method of reducing gastric acid secretion comprising  
administering to a mammal an effective amount of a sulfonyl prodrug of a  
proton pump inhibitor in a composition suitable for said administration,  
20 provided said composition does not comprise a phosphate buffer.
17. The method of claim 16 wherein the proton pump inhibitor is  
lansoprazole.
18. The method of claim 16 wherein the proton pump inhibitor is  
esomeprazole.
- 25 19. The method of claim 16 wherein the proton pump inhibitor is  
omeprazole.
20. The method of claim 16 wherein the proton pump inhibitor is  
pantoprazole.
21. The method of claim 16 wherein the proton pump inhibitor is  
30 rabeprazole.

22. The method of claim 16 wherein said biological leaving group comprises a phenylsulfonyl group, wherein the sulfur atom is directly bonded to the nitrogen atom of the benzimidazole moiety.

23. The method of claim 16 comprising



or a pharmaceutically acceptable salt thereof.

24. The method of claim 16 wherein said prodrug is administered in a dosage form or a composition which does not comprise a polyvalent anion having a molecular mass of 102 or less.

10 25. The method of claim 16 wherein said prodrug is administered in a dosage form or a composition which does not comprise a buffer.

26. The method of claim 16 wherein said prodrug is administered in a dosage form or a composition which does not comprise more than 0.05 moles of a polyvalent anion for every 1 mole of said prodrug, wherein said polyvalent  
15 anion has an aqueous solubility of 0.15 M or greater.

27. The method of claim 16 wherein said prodrug is administered in a dosage form or a composition which does not comprise a polyvalent anion having an aqueous solubility of 0.2 M or greater.

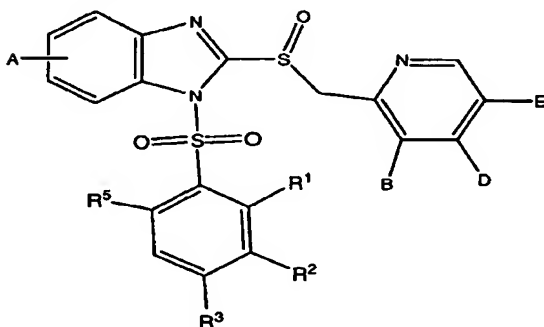
28. The method of claim 16 wherein said prodrug is administered in a dosage form or a composition which does not comprise a polyvalent anion having an aqueous solubility of 0.02 M or greater.  
20

29. The method of claim 16 wherein said prodrug is administered in a dosage form or a composition which does not comprise an anion having an aqueous solubility of 0.15 M or greater and a molecular mass of 120 or less.

25 30. The method of claim 19 wherein said prodrug is administered in a dosage form or a composition which does not comprise an anion having an aqueous solubility of 0.015 M or greater and a molecular mass of 120 or less.

31. A pharmaceutical product comprising  
a composition comprising sulfonamide prodrug of a proton pump inhibitor, and  
a package for dispensing or storing said prodrug,  
wherein said composition does not comprise an anionic buffer.

5 32. The product of claim 25 comprising



or a pharmaceutically acceptable salt thereof  
wherein

A is H, OCH<sub>3</sub>, or OCHF<sub>2</sub>;

10 B is CH<sub>3</sub> or OCH<sub>3</sub>;

D is OCH<sub>3</sub>, OCH<sub>2</sub>CF<sub>3</sub>, or O(CH<sub>2</sub>)<sub>3</sub>OCH<sub>3</sub>;

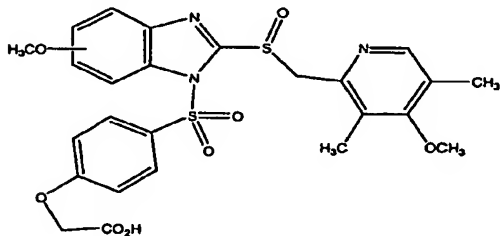
E is H or CH<sub>3</sub>;

R<sup>1</sup>, R<sup>2</sup>, R<sup>3</sup>, and R<sup>5</sup> are independently H, CH<sub>3</sub>, CO<sub>2</sub>H, CH<sub>2</sub>CO<sub>2</sub>H, (CH<sub>2</sub>)<sub>2</sub>CO<sub>2</sub>H,  
CH(CH<sub>3</sub>)<sub>2</sub>, OCH<sub>2</sub>C(CH<sub>3</sub>)<sub>2</sub>CO<sub>2</sub>H, OCH<sub>2</sub>CO<sub>2</sub>CH<sub>3</sub>, OCH<sub>2</sub>CO<sub>2</sub>H, OCH<sub>2</sub>CO<sub>2</sub>NH<sub>2</sub>,

15 OCH<sub>2</sub>CONH<sub>2</sub>(CH<sub>2</sub>)<sub>5</sub>CO<sub>2</sub>CH<sub>3</sub>, or OCH<sub>3</sub>.

33. The product of claim 32 wherein R<sup>1</sup>, R<sup>2</sup>, R<sup>3</sup>, and R<sup>5</sup> are independently H,  
CH<sub>3</sub>, CO<sub>2</sub>H, CH<sub>2</sub>CO<sub>2</sub>H, (CH<sub>2</sub>)<sub>2</sub>CO<sub>2</sub>H, OCH<sub>2</sub>CO<sub>2</sub>CH<sub>3</sub>, OCH<sub>2</sub>CO<sub>2</sub>H,  
OCH<sub>2</sub>CONH<sub>2</sub>(CH<sub>2</sub>)<sub>5</sub>CO<sub>2</sub>CH<sub>3</sub>, or OCH<sub>3</sub>.

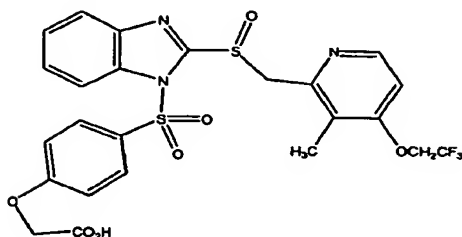
34. The product of claim 31 comprising



20

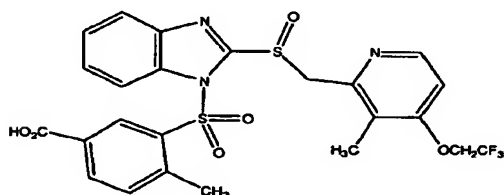
or a pharmaceutically acceptable salt thereof.

35. The product of claim 31 comprising



or a pharmaceutically acceptable salt thereof.

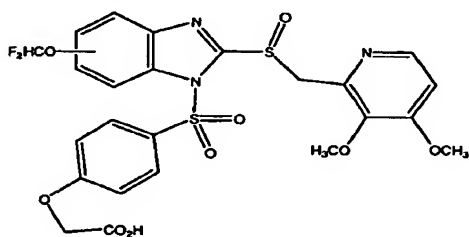
36. The product of claim 31 comprising



5

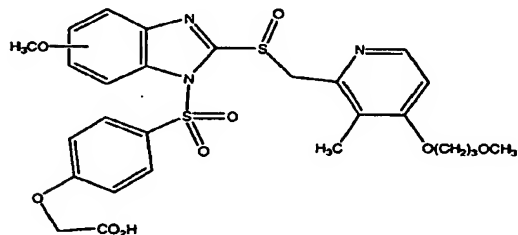
or a pharmaceutically acceptable salt thereof.

37. The product of claim 31 comprising



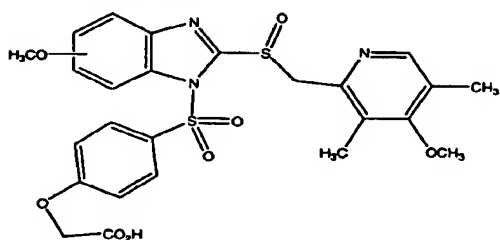
or a pharmaceutically acceptable salt thereof.

10 38. The product of claim 31 comprising



or a pharmaceutically acceptable salt thereof.

39. The product of claim 31 comprising



or a pharmaceutically acceptable salt thereof.

40. The dosage form of claim 1 comprising a buffer which is not anionic.  
5 41. The dosage form of claim 1 which is a liquid.  
42. The dosage form of claim 1 which is a solution.  
43. The dosage form of claim 1 which is a suspension or an emulsion.

ABSTRACT

Disclosed herein are methods, compositions, and dosage forms related to prodrugs of a proton pump inhibitors wherein said compositions and dosage  
5 forms do not comprise a salt of phosphoric acid. Principles related to the use of various anions and buffers in relation to these prodrugs are also disclosed.

**Figure 1**

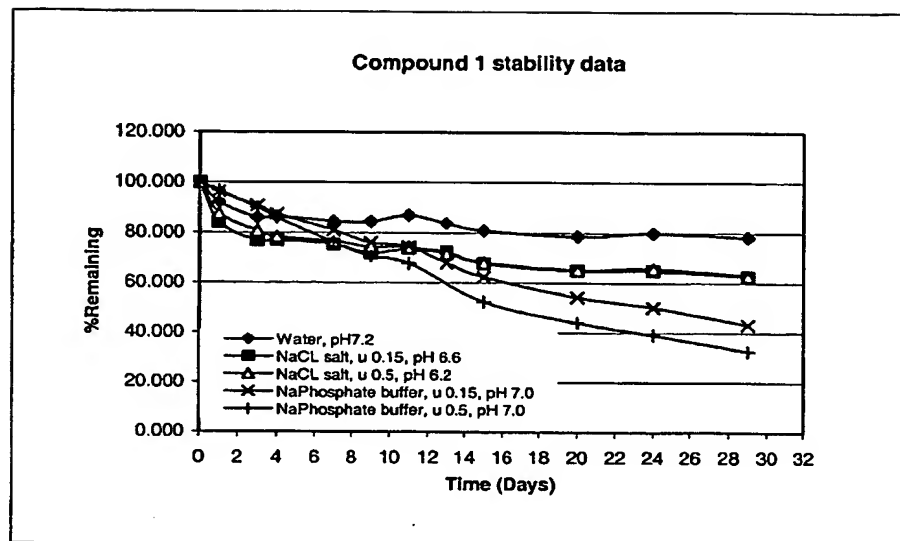


Figure 2

